

**K980213 DI-2000**Mar 17, 1998  
67 days to decisionK980213 · Product code: **LMA** · Radiology  
Source: <https://www.510kdatabase.net/k980213/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Digitizer, Image, Radiological (LMA)
Date received	Jan 9, 1998
Decision date	Mar 17, 1998
Days to decision	67 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lumisys, Inc.</b>
Location	San Leandro, CA, US
Contact	TRINDY LEFORGE
510(k) history	7 submissions · 7 cleared · 1990-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k980213/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 19, 2026